## DEPARTMENT OF HEALTH AND HUMAN SERVICES

Display Date\_\_\_

Publication Date

Certifier

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Food and Drug Administration

21 CFR Part 522

Implantation or Injectable Dosage Form New Animal Drugs; Change of

**Sponsor; Correction** 

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Final rule; correction.

**SUMMARY:** The Food and Drug Administration (FDA) is correcting a final rule that appeared in the **Federal Register** of June 10, 2003 (68 FR 34533). The document amended the animal drug regulations to reflect a change of sponsor for two approved new animal drug applications (NADAs) from Anthony Products Co. to Cross Vetpharm Group Ltd. The document was published with an error. This document corrects that error.

**FOR FURTHER INFORMATION CONTACT:** Joyce A. Strong, Office of Policy (HF-27), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–7010.

**SUPPLEMENTARY INFORMATION:** In the FR Doc. 03–14547, appearing on page 34533 in the **Federal Register** of Tuesday, June 10, 2003, the following correction is made:

## § 522.1696b [Corrected]

■ 1. On page 34534, in the second column, the last line in the amendatory language for § 522.1696b *Penicillin G procaine aqueous suspension* is corrected to read "010515, 053501, 059130, and 61623".

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NADA 129 - 83/

NCR

Dated: 7/7/03

July 7, 2003.

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[FR Doc. 03-????? Filed ??-??-03; 8:45 am]

BILLING CODE 4160-01-S

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